

Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

Ciproxin 500 mg Film-coated Tablets.

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each tablet contains 500 mg ciprofloxacin (as hydrochloride).

For full list of excipients, see section 6.1.

3 PHARMACEUTICAL FORM

Film-coated tablet.

Product imported from the Netherlands, Greece and Hungary:

White, oblong tablets with 'Bayer' impressed on one side and 'CIP', a breakline and '500' on the other.

4 CLINICAL PARTICULARS

4.1 Therapeutic Indications

Ciprofloxacin is indicated for the treatment of the following infections caused by sensitive bacteria:

Adults:

Respiratory tract infections: e.g. lobar and bronchopneumonia, acute and chronic bronchitis, acute exacerbation of cystic fibrosis, bronchiectasis, empysema. Ciprofloxacin is not recommended as first-line therapy for the treatment of pneumococcal pneumonia. In circumstances where a physician considers it appropriate to use ciprofloxacin in patients with pneumococcal pneumonia, a dose of 750mg twice daily should provide adequate cover in the majority of cases (*see Dosage and Administration section*). Ciprofloxacin may be used for treating Gram-negative pneumonia.

Urinary tract infections: e.g. uncomplicated and complicated urethritis, cystitis, pyelonephritis, prostatitis, epididymitis.

Gastro-intestinal infections: e.g. enteric fever, infective diarrhoea.

Gonorrhoea: including urethral, rectal and pharyngeal gonorrhoea caused by β -lactamase producing organisms or organisms moderately sensitive to penicillin.

Children: For the treatment of acute pulmonary exacerbation of cystic fibrosis associated with *P. aeruginosa* infection in paediatric patients aged 5-17 years.

Inhalation Anthrax in Adults and Children: To reduce the incidence or progression of disease following confirmation or suspected exposure to aerosolized *Bacillus anthracis*.

4.2 Posology and method of administration

General dosage recommendations: The dosage of Ciproxin tablets is determined by the severity and type of infection, the sensitivity of the causative organism(s) and the age, weight and renal function of the patient. Ciproxin tablets should be swallowed whole with an adequate amount of liquid.

If ciproxin Tablets are taken on an empty stomach, the active substance is absorbed more rapidly. In this case, the tablets should not be taken concurrently with dairy products or with mineral fortified drinks alone (e.g. milk, yoghurt, calcium fortified orange juice). However, a normal diet that will contain small amounts of calcium, does not significantly affect ciprofloxacin absorption.

Adults: The dosage range for adults is 100-750mg twice daily. The following dosages for specific types of infection are recommended:

Recommended Adult Dosage:

Indication	Dose
Gonorrhoea: 100mg b.d. or 250mg b.d.**	250mg single dose
Upper and lower urinary tract infections (depending on severity):	250-500mg b.d.
Upper and lower respiratory tract infections (depending on severity):	250-750mg b.d.
Pneumococcal pneumonia (second-line):	750mg b.d.
Cystic fibrosis patients with pseudomonal lower RTI*:	750mg b.d.
Other infections as detailed under 4.1:	500-750mg b.d.
Severe infections, particularly due to Pseudomonas, staphylococci and streptococci:	750mg b.d.
inhalation anthrax:	500mg b.d

* As the pharmacokinetics of ciprofloxacin remain unchanged in patients with cystic fibrosis, the low bodyweight of these patients should be taken into consideration when determining dosage.

** Both doses are equally effective in treating acute, uncomplicated cystitis.

Impaired Renal Function:

Dosage adjustments are not usually required, except in patients with severe renal impairment (serum creatinine >265 micromole/l or creatinine clearance <20ml/minute). If adjustment is necessary, this may be achieved by reducing the total daily dose by half, although monitoring of drug serum levels provides the most reliable basis for dose adjustment. Dialysis reduces serum levels of ciprofloxacin.

Elderly:

Although higher ciprofloxacin serum levels are found in the elderly, no adjustment of dosage is necessary.

Adolescents and children:

As with other drugs in its class, ciprofloxacin has been shown to cause arthropathy in weight-bearing joints of immature animals. Although a analysis of available safety data from ciprofloxacin use in patients less than 18 years of age, the majority of whom had cystic fibrosis, did not disclose any evidence of drug-related cartilage or articular damage, its use in the paediatric population is generally not recommended.

Clinical and pharmacokinetic data support the use of ciprofloxacin in paediatric cystic fibrosis patients (aged 5-17 years) with acute pulmonary exacerbation associated with P.aeruginosa infection, at a dose of 20mg/kg orally twice daily (maximum daily dose 1500mg).

For the indication of inhalation anthrax, the risk benefit assessment indicates that administration of ciprofloxacin to paediatric patients at a dose of 15 mg/kg orally twice daily (maximum daily dose of 1000mg) is appropriate.

For indications other than treatment of pulmonary exacerbations in cystic fibrosis and inhalation anthrax, ciprofloxacin may be used in children and adolescents where the benefit is considered to outweigh the potential risks.

Dosing in children with impaired renal and/or hepatic function has not been studied.

Duration of Treatment:

The duration of treatment depends upon the severity of infection, clinical response and bacteriological findings.

In acute, uncomplicated cystitis the treatment period is three days with Ciproxin 100mg Tablets or Ciproxin 250mg

Tablets.

In other acute infections the usual treatment period is 5 to 7 days with Ciproin infusion or 5 to 10 days with Ciproxin Tablets. Generally, treatment should be continued for at least three days after signs and symptoms of the infection have disappeared.

Prolonged treatment or use in chronic conditions should only be initiated under consultant direction with regular surveillance.

For acute pulmonary exacerbation of cystic fibrosis associated with *P.aeruginosa* infection in paediatric patients (aged 5-17 years), the duration of treatment is 10-14 days.

For inhalation anthrax, drug administration should begin as soon as possible after confirmed or suspected exposure and should be continued for 60 days.

4.3 Contraindications

Ciprofloxacin is contraindicated in patients who have shown hypersensitivity to ciprofloxacin or other quinolone anti-infectives or who have a history of quinolone-induced tendon disorder.

Ciprofloxacin is also contraindicated in children and growing adolescents unless epiphyseal closures of long bones have occurred or except where the benefits of treatment exceed the risks.

See 4.2, Adolescents and Children

4.4 Special warnings and precautions for use

In the event of hypersensitivity, therapy should be discontinued.

Ciprofloxacin should be used with caution in epileptics and patients with a history of CNS disorders and only if the benefits of treatment are considered to outweigh the risk of possible CNS side-effects, depression or psychoses lead to self-endangering behaviour (*see also Section 4.8*).

Crystalluria related to the use of ciprofloxacin has been reported. Patients receiving ciprofloxacin should be well hydrated and excessive alkalinity of the urine should be avoided.

Patients with a family history of or actual defects in glucose-6-phosphate dehydrogenase activity are prone to haemolytic reactions with quinolones, and so ciprofloxacin should be used with caution in these patients.

Ciprofloxacin is not recommended as first-line therapy for the treatment of pneumococcal pneumonia. *Streptococcus pneumoniae* is the most frequent pathogen responsible for community acquired pneumonia.

Tendon inflammation and rupture may occur with quinolone antibiotics. Such reactions have been observed particularly in older patients and in those treated concurrently with corticosteroids. At the first sign of pain or inflammation, patients should discontinue ciprofloxacin and rest the affected limbs.

Toxicological studies have shown that administration of oxyquinolone antibacterial agents at doses higher than the therapeutic range can produce erosion of the cartilage in weight-bearing joints in immature animals of some species. No such lesions have been shown to occur in man to date. This product should not be prescribed for children or those in whom bone growth is continuing, with the exception of paediatric cystic fibrosis patients, unless the benefit of short-term use is regarded as exceeding the risk.

Patients with pre-existent significant renal or hepatic disorders should be carefully monitored to detect any deterioration in function. It should only be administered with great caution to persons with renal insufficiency, or severe dehydration.

There is a risk of pseudomembranous colitis with broad-spectrum antibiotics. It is important to consider this in patients suffering from severe, persistent diarrhoea. With ciprofloxacin this effect has been reported rarely. If

pseudomembranous colitis is suspected treatment with ciprofloxacin should be stopped and appropriate treatment given (e.g. oral vancomycin).

As with other quinolones, patients should avoid prolonged exposure to strong sunlight or UV radiation during treatment.

Laboratory tests may give abnormal findings if performed whilst patients are receiving ciprofloxacin, e.g. increased alkaline phosphatase; increases in liver function tests, e.g. transaminases and cholestatic jaundice, especially in patients with previous liver damage.

Eradication of infection due to *Pseudomonas* in persons with cystic fibrosis only occurs in a minority of cases, particularly after repeat courses of treatment with ciprofloxacin. Cyclical or alternating antibacterial therapies may help reduce the number of resistant strains.

4.5 Interaction with other medicinal products and other forms of interaction

Increased plasma levels of theophylline have been observed following concurrent administration with ciprofloxacin. It is recommended that the dose of theophylline should be reduced and plasma levels of theophylline monitored. Where monitoring of plasma levels is not possible, the use of ciprofloxacin should be avoided in patients receiving theophylline. Particular caution is advised in those patients with convulsive disorders.

Phenytoin levels may be altered when Ciproxin is used concomitantly.

Ciproxin Tablets should not be administered within 4 multivalent cationic drugs and mineral supplements (e.g. calcium, magnesium, aluminium or iron), sucralfate or antacids and highly buffered drugs (e.g. anti retrovirals) as interference with absorption may occur. When appropriate, patients should be advised not to self-medicate with preparations containing these compounds during therapy with ciprofloxacin. This reaction does not apply to the class of H₂ receptor blocker drugs.

The concurrent administration of dairy products or fortified drinks alone (e.g. milk, yoghurt, calcium, fortified orange juice) and ciprofloxacin should be avoided because absorption of ciprofloxacin may be reduced. However a normal diet, that will contain small amounts of calcium, does not significantly affect ciprofloxacin absorption.

Prolongation of clotting time has been reported during concomitant administration of ciprofloxacin and oral anti-coagulants.

Ciprofloxacin may interfere with estimations of urinary 17-ketosteroids, or vanillylmandelic acid.

Animal data have shown that high doses of quinolones in combination with some non-steroidal anti-inflammatory drugs, (e.g. fenbufen, but not acetylsalicylic acid) can lead to convulsions.

Transient increases in serum creatinine have been seen following concomitant administration of ciprofloxacin and cyclosporin. Therefore, monitoring of serum creatinine levels is advisable.

Concomitant use with some phenylpropionic acid-derived non-steroidal anti-inflammatory drugs may lead to toxicity possible because of renal effects.

The simultaneous administration of quinolones and glibenclamide can on occasion potentiate the effect of glibenclamide resulting in hypoglycaemia.

Renal tubular transport of methotrexate may be inhibited by concomitant administration of ciprofloxacin potentially leading to increased plasma levels of methotrexate. This may increase the risk of methotrexate associated toxic reactions. Therefore, patients receiving methotrexate therapy should be carefully monitored when concomitant ciprofloxacin therapy is indicated.

Concomitant use with probenecid reduces the renal clearance of ciprofloxacin, resulting in increased quinolone plasma levels.

The use of metoclopramide with ciprofloxacin may accelerate the absorption of ciprofloxacin.

4.6 Fertility, pregnancy and lactation

Ciprofloxacin should not be used during pregnancy, or in women at risk of pregnancy nor during lactation.

Reproduction studies performed in mice, rats and rabbits using parenteral and oral administration did not reveal any evidence of teratogenicity, impairment of fertility or impairment of peri-/post-natal development. However, as with other quinolones, ciprofloxacin has been shown to cause arthropathy in immature animals, and therefore its use during pregnancy or in women capable of child-bearing is not recommended. Studies have indicated that ciprofloxacin is secreted in breast milk. Administration to nursing mothers is thus not recommended.

4.7 Effects on ability to drive and use machines

Ciprofloxacin could result in impairment of the patient's ability to drive or operate machinery, particularly in conjunction with alcohol.

4.8 Undesirable effects

Ciprofloxacin is generally well tolerated. The most frequently reported adverse reactions are nausea, diarrhoea and rash.

The following adverse reactions have been observed:

Hypersensitivity/skin, e.g. rash, pruritus, urticaria, photosensitivity, drug-induced fever, anaphylactic/anaphylactoid reactions including angioedema and dyspnoea. Rarely, erythema nodosum and erythema multiforme. Very rarely, petechiae, haemorrhagic bullae, vasculitis, serum sickness-like reaction, fixed drug reactions, Stevens-Johnson Syndrome and Lyells Syndrome. Treatment with ciprofloxacin should be discontinued if any of the above occur upon first administration.

General, e.g. moniliasis, asthenia, hyperglycaemia and abnormal gait, pain, pain in extremities, back pain, chest pain. Thrombophlebitis.

CNS, e.g. headache, restlessness, depression, dizziness, tremor, convulsions, confusion, hallucinations, somnolence, sleep disorders, insomnia. Very rarely, migraine, hyperesthesia, ataxia, hypertonia, twitching or and anxiety states. Isolated cases of ciprofloxacin-induced psychoses have been reported, which may progress to self-endangering behaviour. There are isolated reports of intracranial hypertension associated with quinolone therapy. Paraesthesia has been reported.

Gastro-intestinal, e.g. nausea, diarrhoea, vomiting, dyspepsia, abdominal pain, anorexia, flatulence, dysphagia. Rarely, pancreatitis or pseudomembranous colitis.

Cardiovascular, e.g. tachycardia, oedema, fainting, hot flushes, hypotension and sweating, 'Very rare prolonged QTc interval and ventricular arrhythmia (including torsade de pointes).

Effects on haematological parameters, e.g. anaemia, eosinophilia, increases or decreases in white cell and/or platelet count, altered prothrombin levels, and, very rarely, haemolytic anaemia, agranulocytosis or pancytopenia or bone marrow depression. Pancytopenia and bone marrow depression may be potentially life threatening.

Hepatic, e.g. transient increases in liver enzymes or serum bilirubin (particularly in patients with previous liver damage), hepatitis, jaundice/cholestasis and major liver disorders including hepatic necrosis, which may rarely progress to hepatic failure.

Renal, e.g. transient increases in blood urea or serum creatinine, renal failure, crystalluria, haematuria, nephritis.

Musculoskeletal, e.g. reversible arthralgia, joint swelling and myalgia. Rarely, tenosynovitis. Tendon inflammation (predominantly of the Achilles tendon) has been reported which may lead to tendon rupture. Treatment should be discontinued immediately if these symptoms occur.

Rarely, exacerbation of the symptoms of myasthenia gravis has been reported.

Special senses, e.g. very rarely, visual disturbances including diplopia and colour disturbances, impaired taste and smell usually reversible upon discontinuation of treatment, tinnitus, transient impairment of hearing particularly at high frequencies.

Abnormal laboratory findings, e.g. increased alkaline phosphatase, amylase and lipase, increases in liver function tests, e.g transaminases, and cholestatic jaundice, especially in patients with previous liver damage.

4.9 Overdose

Based on the limited information available in two cases of ingestion of over 18g of ciprofloxacin, reversible renal toxicity has occurred. Therefore, apart from routine emergency measures, it is recommended to monitor renal function, including urinary pH and acidify, if required, to prevent crystalluria.

Patients must be kept well hydrated and, in the case of renal damage resulting in prolonged oliguria, dialysis should be initiated.

Calcium or magnesium antacids may be administered as soon as possible after ingestion of Ciproxin tablets in order to reduce the absorption of ciprofloxacin.

Serum levels of ciprofloxacin are reduced by dialysis.

5 PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties

ATC Code J01 MA 02

Ciprofloxacin is a synthetic 4-quinolone derivative, with bactericidal activity. It acts via inhibition of bacterial DNA gyrase, ultimately resulting in interference with DNA function. Ciprofloxacin is highly active against a wide range of Gram-positive and Gram-negative organisms and has shown activity against some anaerobes, Chlamydia spp.and Mycoplasma spp.. Killing curves demonstrate the rapid bactericidal effect against sensitive organisms and it is often found that minimum bactericidal concentrations are in the range of minimum inhibitory concentrations

Ciprofloxacin has been shown to have no activity against *Treponema pallidum* and *Urealasma urealyticum*, *Nocardia asteroides* and *Enterococcus faecium* are resistant.

Breakpoints

S ≤ 1µg/ml, R ≥ 4µg/ml

Susceptibility:

The prevalence of resistance may vary geographically and with time for selected species and local area information on resistance is desirable, particularly when treating severe infections. This information gives only an approximate guidance on whether micro-organisms will be susceptible to ciprofloxacin or not.

Organism	Prevalence of resistance
Sensitive:	
Gram-positive bacteria	
Corynebacterium diphtheriae	0%
Corynebacterium spp.	-
Staphylococcus aureus (methicillin sensitive)	0-14%

Staphylococcus aureus (methicillin aresistant)	48-90%
Streptococcus agalactiae	0-17%
Bacillus anthracis	-

Gram-negative bacteria	
Acinetobacter baumannii	6-93%
Acinetobacter spp.	14-70%
Aeromonas hydrophilia	0%
Aeromonas spp.	-
Bordetella pertussis	0%
Brucella melitensis	0%
Campylobacter jejuni/coli	0.82%
Campylobacter spp.	0%
Citrobacter freundii	0-4%
Citrobacter spp.	0%
Edwardsiella tarda	0%
Enterobacter aerogenes	0%
Enterobacter cloacae	0-3%
Enterobacter spp.	3-13%
Escherichia coli	2-7%
Escherichia coli, EHEC and EPEC	-
Haemphilus influenzae	0-1%
Haemphilus influenzae (β-lactam negative)	0%
Haemphilus influenzae (β-lactam postive)	0%
Haemphilus parainfluenzae	0%
Hafnia alvei	0%
Klebsiella oxytoca	0%
Klebsiella pneumoniae	2-5.8%
Klebsiella spp.	2-21%
Legionella pneumophila	0%
Legionella spp.	0%
Moraxella catarrhalis	0%
Morganella morganii	1-2%
Neisseria gonorrhoeae	0%
Neisseria gonorrhoeae, β-lactamase	0%
Neisseria gonorrhoeae, β-lactamase positive	0%
Neisseria meningitidis	0%
Neisseria meningitidis, β-lactamase negative	0%
Plesiomonas shigelloides	0%
Proteus mirabillis	0-10%
Proteus vulgaris	4%
Providence rettgeri	-
Providence spp.	4%
Providence stuartii	-
Pseudomonas aeruginosa	1-28%
Salmonella spp.	0%
Salmonella typhi	0-2%
Serratia liquefaciens	-
Serratia marcescens	23%
Serratia spp.	0-21%
Shigella spp.	0%

Vibrio cholerae	0%
Vibrio parahaemolyticus	0%
Vibrio spp.	0%
Yershinia enterocolitica	0%

Anaerobes	
Bacteroides ureolyticus	0%
Clostridium perfringens	-
Peptococcus spp.	0%
Pepotostreptococcus spp.	-
Peptostreptococcus magnus	0%
Veillonella parvula	0%
Other pathogens	
Chlamydia spp.	-
Heilicobacter pylori	-
Mycobacterium fortuitum	0%
Mycobacterium tuberculosis	0%
Mycoplasma hominis	16%
Intermediate	
Gram-positive aerobes	
Enterococci	5%
Enterococcus faecalis	9-34%
Staphylococcus epidermis, methicillin sensitive	10-16%
Staphylococcus epidermis, methicillin resistant	26-56%
Staphylococcus haemolyticus	-
Staphylococcus haemolyticus, methicillin sensitive	8%
Staphylococcus haemolyticus, methicillin resistant	73%
Streptococcus anginosus	9%
Streptococcus bovis	-
Streptococcus milleri	5%
Streptococcus mitis	-
Streptococcus pneumoniae, penicillin sensitive	0-1%
Streptococcus pneumoniae, penicillin intermediate	-
Streptococcus pneumoniae, penicillin intermediate and resistant	2.8%
Streptococcus pneumoniae, penicillin resistant	-
Streptococcu pyogenes	0-28%
Streptococcus, viridans group	-
Streptococcus viridans, penicillin sensitive	-
Streptococcus viridans, penicillin resistant	-
Streptococcu, β-haemolytic groups A,C and G	0%
Gram-negative aerobes	
Alcaligenes spp.	-
Listeria monocytogenes	0%
Listeria spp.	0%
Anaerobes	

Fusobacterium spp.	-
Gardnerella vaginalis	0%
Preotella spp.	-

Other pathogens	
Ureaplasma urealyticum	11%
Resistant	
Gram-postive aerobes	
Enterococcus faecium	-
Stenotrophomonas maltophilla	94%
Streptococcus samguis	-
Gram-negative aerobes	
Flavobacterium meningosepticum	-
Nocardia asteroides	-
Anaerobes	
Bacteroides fragilis	-
Bacteroides thetaiotaomicron	-
Clostridium difficile	-

Reistance

Plasmid-related transfer of resistance has not been observed with ciprofloxacin and the overall frequency of development of resistance is low (10-9 - 10-7). Cross-resistance to penicillins, cephalosporins, aminoglycosides and tetracyclines has not been observed and organisms resistant to these antibiotics are generally sensitive to ciprofloxacin. Ciprofloxacin is also suitable for use in combination with these antibiotics, and additive behaviour is usually observed.

5.2 Pharmacokinetic properties

Absorption of oral doses of ciprofloxacin tablet formulation occurs rapidly, mainly from the small intestine, the half-life of absorption being 2-15 minutes. Plasma levels are dose-related and peak 0.5-2.0 hours after dosing. The AUC also increases dose proportionately after administration of both single and repeated oral (tablet) and intravenous doses. The absolute bioavailability is reported to be 52-83% and ciprofloxacin is subject to only slight first pass metabolism. The oral bioavailability is approximately 70-80%.

The intake of food at the same time as administration of oral ciprofloxacin has a marginal but clinically not relevant effect on the pharmacokinetic parameters Cmax and AUC. No specific recommendations are necessary with regard to time of administration of oral ciprofloxacin relative to food intake. Distribution of ciprofloxacin within tissues is wide and the volume of distribution high, though slightly lower in the elderly. Protein binding is low (between 19-40%).

Only 10-20% of a single oral or intravenous dose is eliminated as metabolites (which exhibit lower activity than the parent drug). Four different antimicrobially active metabolites have been reported, desethyleneciprofloxacin (M1), sulphociprofloxacin (M2), oxaciprofloxacin (M3) and formylciprofloxacin (M4). M2 and M3 account for one third each of metabolised substance and M1 is found in small amounts (1.3-2.6% of the dose). M4 has been found in very small quantities (<0.1% of the dose). M1-M3 have antimicrobial activity comparable to nalidixic acid and M4 found in the smallest quantity has antimicrobial activity similar to that of norfloxacin.

Elimination of ciprofloxacin and its metabolites occurs rapidly, primarily by the kidney. After single oral and intravenous doses of ciprofloxacin, 55% and 75% respectively are eliminated by the kidney and 39% and 14% in the faeces within 5 days. Renal elimination takes place mainly during the first 12 hours after dosing and renal clearance levels suggest that active secretion by the renal tubules occurs in addition to normal glomerular filtration. Renal

clearance is between 0.18 – 0.3 l/h.kg and total body clearance between 0.48 – 0.60 l/h.kg. Approximately 1% of a ciprofloxacin dose is excreted via the biliary route. The elimination kinetics are linear and after repeated dosing at 12 hourly intervals, no further accumulation is detected after the distribution equilibrium is attained (at 4-5 half-lives). The elimination half-life of unchanged ciprofloxacin over a period of 24-48 hours post-dose is 3.1-5.1 hours.

Some studies carried out with ciprofloxacin in severely renally impaired patients (serum creatinine >265 micromole/l or creatinine clearance <20ml/minute) demonstrated either a doubling of the elimination half-life, or fluctuations in half-life in comparison with healthy volunteers, whereas other studies showed no significant correlation between elimination half-life and creatinine clearance. However, it is recommended that in severely renally impaired patients, the total daily dose should be reduced by half, although monitoring of drug serum levels provides the most reliable basis for dose adjustment as necessary.

Results of pharmacokinetic studies in paediatric cystic fibrosis patients have shown dosages of 20mg/kg orally twice daily or 10mg/kg iv three times daily are recommended to achieve plasma concentration/time profiles comparable to those achieved in the adult population at the currently recommended dosage regimen.

Inhalation anthrax: Ciprofloxacin serum concentrations achieved in humans serve as a surrogate endpoint reasonably likely to predict clinical benefit and provide the basis for the recommended doses.

5.3 Preclinical safety data

Following extensive oral and intravenous toxicology testing with ciprofloxacin, only two findings which may be considered relevant to the use of ciprofloxacin in man were observed. Crystalluria was noted in those species of animals which had a normally alkaline urine. Kidney damage without the presence of crystalluria was not observed. This effect is considered a secondary inflammatory foreign-body reaction, due to the precipitation of a crystalline complex of ciprofloxacin, magnesium and protein in the distal tubule system of the kidneys. This is considered not to be a problem in man, because the urine is normally acidic. However, to avoid the occurrence of crystalluria, patients should be well hydrated and excessive alkalinity of the urine avoided.

As with other quinolones, damage to the weight-bearing joints of only juvenile rats and dogs treated with ciprofloxacin was noted in repeat dose toxicity testing. This was more noticeable in the dog. Although analysis of available safety data from ciprofloxacin use in paediatric patients did not disclose any evidence of drug related cartilage or articular damage, the use of ciprofloxacin in children and growing adolescents is generally not recommended, unless the benefits are considered to outweigh the potential risks (with the exception of treatment of cystic fibrosis). Additionally, because of the potential of arthropathy, the use of ciprofloxacin during pregnancy and lactation is not recommended.

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Maize starch
Microcrystalline cellulose
Crospovidone
Colloidal anhydrous silica
Magnesium stearate
Hypromellose
Macrogol 4000
Titanium dioxide (E 171)

6.2 Incompatibilities

Not applicable.

6.3 Shelf life

The shelf-life expiry date of this product shall be the date shown on the container and outer package of the product on the market in the country of origin.

6.4 Special precautions for storage

Do not store above 30°C.
Store in the original package.

6.5 Nature and contents of container

Blister packs of 10 and 20 tablets.

6.6 Special precautions for disposal of a used medicinal product or waste materials derived from such medicinal product and other handling of the product

No special requirements.

7 PARALLEL PRODUCT AUTHORISATION HOLDER

PCO Manufacturing
Unit 10, Ashbourne Business Park
Rath
Ashbourne
Co. Meath

8 PARALLEL PRODUCT AUTHORISATION NUMBER

PPA 0465/082/002

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 22 March 2002

Date of last renewal: 22 March 2007

10 DATE OF REVISION OF THE TEXT

April 2011